

BA

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
17 March 2005 (17.03.2005)

PCT

(10) International Publication Number
WO 2005/023264 A1

- (51) International Patent Classification⁷: **A61K 31/497**, 31/663, 31/675, 38/09, A61P 35/00
- (74) Agent: GLOBAL INTELLECTUAL PROPERTY; AstraZeneca AB, S-151 85 Södertälje (SE).
- (21) International Application Number: PCT/GB2004/003733
- (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (22) International Filing Date: 2 September 2004 (02.09.2004)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data: 0320806.3 5 September 2003 (05.09.2003) GB
- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).
- (71) Applicant (for all designated States except MG, US): ASTRAZENECA AB [SE/SE]; S-151 85 Södertälje (SE).
- (71) Applicant (for MG only): ASTRAZENECA UK LIMITED [GB/GB]; 15 Stanhope Gate, London Greater London W1K 1LN (GB).
- (72) Inventor; and
- (75) Inventor/Applicant (for US only): GALLAGHER, Nell [IE/GB]; AstraZeneca R & D Alderley, Alderley Park, Macclesfield Cheshire SK10 4TG (GB).
- Published:
— with international search report
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



WO 2005/023264 A1

(54) Title: COMBINATION COMPRISING N-(3-METHOXY-5-METHYLPYRAZIN-2-YL)-2-(4-[1,3,4-OXADIAZOL-2-YL]PHENYL)PYRIDINE-3-SULPHONAMIDE AND AN LHRH ANALOGUE AND/OR A BISPHOSPHONATE

(57) Abstract: A combination, comprising N-(3-methoxy-5-methylpyrazin-2-yl)-2-(4-[1,3,4-oxadiazol-2-yl]phenyl)pyridine-3-sulphonamide, or a pharmaceutically acceptable salt thereof, and an LHRH analogue and / or a bisphosphonate is described.

COMBINATION COMPRISING N- (3-METHOXY-5-METHYLPYRAZIN-2-YL) -2- (4- (1,3,4- OXADIAZOL-2-YL) PHENYL) PYRIDINE-3-SULPHONAMIDE AND AN LHRH ANALOGUE AND/OR A BIPHOSPHONATE

The present invention relates to a combination comprising *N*-(3-methoxy-5-methylpyrazin-2-yl)-2-(4-[1,3,4-oxadiazol-2-yl]phenyl)pyridine-3-sulphonamide, or a
5 pharmaceutically acceptable salt thereof, hereafter "Compound (I)", and a luteinizing hormone releasing hormone analogue (hereafter LHRH analogue). The present invention further relates to a combination comprising Compound (I) and a diphosphonic acid derivative (hereafter bisphosphonate) and a combination comprising Compound (I), an LHRH analogue and a bisphosphonate.

10 These combinations are useful for the treatment or prophylaxis of cancer. The invention also relates to a pharmaceutical composition comprising such combinations and to the use thereof in the manufacture of a medicament for use in the treatment or prophylaxis of cancer, in particular prostate cancer.

Cancer affects an estimated 10 million people worldwide. This figure includes
15 incidence, prevalence and mortality. More than 4.4 million cancer cases are reported from Asia, including 2.5 million cases from Eastern Asia, which has the highest rate of incidence in the world. By comparison, Europe has 2.8 million cases, North America 1.4 million cases, and Africa 627,000 cases.

In the UK and US, for example, more than one in three people will develop cancer at
20 some point in their life. Cancer mortality in the U.S. is estimated to account for about 600,000 a year, about one in every four deaths, second only to heart disease in percent of all deaths, and second to accidents as a cause of death of children 1-14 years of age. The estimated cancer incidence in the U.S. is now about 1,380,000 new cases annually, exclusive of about 900,000 cases of non-melanotic (basal and squamous cell) skin cancer.

25 Cancer is also a major cause of morbidity in the UK with nearly 260,000 new cases (excluding non-melanoma skin cancer) registered in 1997. Cancer is a disease that affects mainly older people, with 65% of cases occurring in those over 65. Since the average life expectancy in the UK has almost doubled since the mid nineteenth century, the population at risk of cancer has grown. Death rates from other causes of death, such as heart disease, have
30 fallen in recent years while deaths from cancer have remained relatively stable. The result is that 1 in 3 people will be diagnosed with cancer during their lifetime and 1 in 4 people will die from cancer. In people under the age of 75, deaths from cancer outnumber deaths from diseases of the circulatory system, including ischaemic heart disease and stroke. In 2000,

- 2 -

there were 151,200 deaths from cancer. Over one fifth (22 per cent) of these were from lung cancer, and a quarter (26 per cent) from cancers of the large bowel, breast and prostate.

Worldwide, the incidence and mortality rates of certain types of cancer (of stomach, breast, prostate, skin, and so on) have wide geographical differences which are attributed to racial, cultural, and especially environmental influences. There are over 200 different types of cancer but the four major types, lung, breast, prostate and colorectal, account for over half of all cases diagnosed in the UK and US. Prostate cancer is the fourth most common malignancy among men worldwide, with an estimated 400,000 new cases diagnosed annually, accounting for 3.9 percent of all new cancer cases.

Current options for treating cancers include surgical resection, external beam radiation therapy and / or systemic chemotherapy. These are partially successful in some forms of cancer, but are not successful in others. There is a clear need for new therapeutic treatments.

Recently, endothelin A receptor antagonists have been identified as potentially of value in the treatment of cancer (Cancer Research, 56, 663-668, February 15th, 1996 and Nature Medicine, Volume 1, Number 9, September 1999, 944-949).

The endothelins are a family of endogenous 21 amino acid peptides comprising three isoforms, endothelin-1, endothelin-2 and endothelin-3. The endothelins are formed by cleavage of the Trp²¹-Val²² bond of their corresponding proendothelins by an endothelin converting enzyme. The endothelins are among the most potent vasoconstrictors known. They exhibit a wide range of other activities including stimulation of cell proliferation and mitogenesis, inhibition of apoptosis, extravasation and chemotaxis, and also interact with a number of other vasoactive agents.

The endothelins are released from a range of tissue and cell sources including vascular endothelium, vascular smooth muscle, kidney, liver, uterus, airways, intestine and leukocytes. Release can be stimulated by hypoxia, shear stress, physical injury and a wide range of hormones and cytokines. Elevated endothelin levels have been found in a number of disease states in man including cancers.

LHRH is a decapeptide that is secreted by the hypothalamus into the hypophyseal portal circulation in response to neural and/or chemical stimuli, causing the biosynthesis and release of luteinizing hormone (LH) and follicle-stimulating hormone (FSH) by the pituitary. LHRH is also known by other names, Gonadotropin releasing hormone (GnRH), gonadoliberin, FSH releasing hormone (FSH RH) and LH/FSH releasing factor (LH/FSH RF).

- 3 -

The role of gonadal androgens in propagating and maintaining the growth of prostate cancer has long been recognised (Huggins C, Hodges CV. *Cancer Res* 1941;1:293). LHRH plays an important role in regulating the action of LH and FSH (by regulation of their levels), and thus has a role in regulating the levels of gonadal steroids in both sexes, including the sex hormones progesterone, oestrogens and androgens. More discussion of LHRH can be found in
5 WO 97/14697, the disclosure of which is incorporated herein by reference.

Medical and surgical castration are commonly used for the treatment of prostate cancer and LHRH analogues are frequently used to induce medical castration for the management of patients with early and advanced prostate cancer. They have proven effective
10 in the treatment of certain conditions which require inhibition of LH/FSH release. In particular, LHRH-based therapies have proven effective in the treatment of endometriosis, uterine fibroids, polycystic ovarian disease, precocious puberty and several gonadal steroid - dependent neoplasia, most notably cancers of the prostate, breast and ovary. LHRH analogues have also been utilized in various assisted fertilization techniques and have been investigated
15 as a potential contraceptive in both men and women. They have also shown possible utility in the treatment of pituitary gonadotrophe adenomas, sleep disorders such as sleep apnea, irritable bowel syndrome, premenstrual syndrome, benign prostatic hyperplasia, hirsutism, as an adjunct to growth hormone therapy in growth hormone deficient children, and in murine models of lupus.

20 Bisphosphonates are diphosphonic acid derivatives that are capable of regulating metal cations content (especially calcium content) in humans. In particular, they have a pronounced regulatory action on the calcium metabolism of warm-blooded animals. They are thus useful in the treatment of diseases connected with content and circulation of these cations particularly the retardation of bone decalcification. Most particularly, they effect a marked
25 inhibition of bone resorption in rats, as can be demonstrated in the experimental procedure described in *Acta Endocrinol.* 78, 613-24 (1975).

The clinical use of bisphosphonates has increased dramatically in the past decade. The most common indication for these compounds is osteoporosis, but their use in osteolytic bone disease has increased rapidly. The FDA approved pamidronate in 1995 for the treatment of
30 normocalcemic patients with myeloma. In 1996 they issued a new approval for patients with osteolytic lesions of metastatic breast cancer prompting the widespread use of pamidronate in patients with tumors that metastasize to bone. Although the effect of bisphosphonates on

- 4 -

osteoclast-mediated bone resorption has been known and well documented for many years, the exact mechanism of that inhibition is still not completely defined.

The present inventors have unexpectedly found that:

- i) the combination use of Compound (I) and LHRH analogues; or
 - 5 ii) the combination use of Compound (I), and bisphosphonates; or
 - iii) the combination use of Compound (I), LHRH analogues and bisphosphonates;
- can have a particular benefit in the treatment of cancer.

Therefore according to the present invention, there is provided a combination, comprising Compound (I), and an LHRH analogue.

10 According to a further aspect of the present invention, there is provided a combination, comprising Compound (I), and a bisphosphonate.

According to a further aspect of the present invention, there is provided a combination, comprising Compound (I), an LHRH analogue and a bisphosphonate.

Herein where the term "LHRH analogue" is used it is to be understood that this refers
15 to any chemical compound, or a pharmaceutically acceptable salt thereof, including small molecules and peptides, which acts as an agonist or antagonist at the LHRH receptor, whether by an interaction with the LHRH binding site or by an allosteric mechanism, i.e. acts at a position on the LHRH receptor different to the LHRH binding site. In one aspect of the invention an "LHRH analogue" refers to an LHRH antagonist or a pharmaceutically
20 acceptable salt thereof. In one aspect of the invention an "LHRH analogue" refers to an LHRH agonist or a pharmaceutically acceptable salt thereof. In a further aspect of the invention an "LHRH analogue" refers to a combination of an LHRH antagonist or a pharmaceutically acceptable salt thereof and an LHRH agonist or a pharmaceutically acceptable salt thereof.

25 A "bisphosphonate" is a compound, or a pharmaceutically acceptable salt thereof, capable of regulating metal cations content (especially calcium content) in humans and is a compound containing two carbon phosphorous bonds. For further explanation of the term "bisphosphonate" the readers attention is drawn to Endocrine Reviews, 1998, 19(1): 80-100, the content of which is incorporated herein by reference.

30 Herein, where the term "combination" is used it is to be understood that this refers to simultaneous, separate or sequential administration. In one aspect of the invention "combination" refers to simultaneous administration. In another aspect of the invention "combination" refers to separate administration. In a further aspect of the invention

- 5 -

“combination” refers to sequential administration. Where the administration is sequential or separate, the delay in administering the second component should not be such as to lose the benefit of the synergistic effect of the combination.

In one aspect, where a compound or a pharmaceutically acceptable salt thereof, is referred to this refers to the compound only. In another aspect this refers to a pharmaceutically acceptable salt of the compound.

Where cancer is referred to, particularly it refers to oesophageal cancer, myeloma, hepatocellular, pancreatic, cervical cancer, ewings tumour, neuroblastoma, kaposi sarcoma, ovarian cancer, breast cancer, colorectal cancer, prostate cancer, bladder cancer, melanoma, lung cancer - non small cell lung cancer (NSCLC), and small cell lung cancer (SCLC), gastric cancer, head and neck cancer, brain cancer, renal cancer, lymphoma and leukaemia. More particularly it refers to prostate cancer. In addition, more particularly it refers to SCLC, NSCLC, colorectal cancer, ovarian cancer and / or breast cancer. In addition, more particularly it refers to SCLC. In addition, more particularly it refers to NSCLC. In addition, more particularly it refers to colorectal cancer. In addition, more particularly it refers to ovarian cancer. In addition, more particularly it refers to breast cancer. Furthermore, more particularly it refers to bladder cancer, oesophageal cancer, gastric cancer, melanoma, cervical cancer and / or renal cancer. In addition it refers to endometrial, liver, stomach, thyroid, rectal and / or brain cancer. In another aspect of the invention, the cancer is not melanoma. In another embodiment of the invention, particularly the cancer is in a metastatic state, and more particularly the cancer produces metastases to the bone. In a further embodiment of the invention, particularly the cancer is in a metastatic state, and more particularly the cancer produces skin metastases. In a further embodiment of the invention, particularly the cancer is in a metastatic state, and more particularly the cancer produces lymphatic metastases. In a further embodiment of the invention, the cancer is in a non-metastatic state.

Where the treatment of cancer is referred to particularly this is the treatment of cancerous tumours expressing endothelin A. This treatment is in terms of one or more of the extent of the response, the response rate, the time to disease progression and the survival rate. It is further expected that the combination use of Compound (I) and particular LHRH analogues will have a beneficial effect in preventing the onset of cancer in warm-blooded animals, such as man.

Particular compounds, or pharmaceutically acceptable salts thereof possessing LHRH analogue activity include Azaline B, A-198401, A-75998, A-76154, A-84861, abarelix,

- 6 -

AN-152, AN-207, Antide, avorelin, cetrorelix, D-21775, D-23487, D-26344, D-63153, D-85108, degarelix, deslorelin, detirelix, FE 200486, ganirelix, gonadimmune, goserelin, histrelin, leuprolide, leuprorelin, metarelin, nafarelin, NBI-42902 (Neurocrine), Org-30850, PH-45 (Pherin Corp), PTL-03001, ramorelix, RWJ-47428-021, SPD-424, surfagon, T-66
 5 (Matrix Therapeutics Ltd), TAK-013, TAK-810, teverelix, triptorelin acetate, triptorelin pamoate, vomeropherin or ZK-157348.

Particular LHRH analogues are peptides or peptide derivatives.

Examples of particular LHRH agonists include, but are not limited to;

- i) buserelin (US Patent 4 024 248)
 10 (pyr)Glu-His-Trp-Ser-Tyr-D-Ser(Bu)⁶-Leu-Arg-Pro-NHCH₂CH₃
- ii) triptorelin (US Patent 4 010 125)
 (pyr)Glu-His-Trp-Ser-Tyr-Trp-Leu-Arg-Pro-Gly-NH₂
- iii) leuprorelin (US Patent 4 005 063)
 (pyr)Glu-His-Trp-Ser-Tyr-D-Leu-Leu-Arg-Pro-NHCH₂CH₃
- 15 iv) goserelin (US Patent 4 100 274)
 (pyr)Glu-His-Trp-Ser-Tyr-D-Ser(Bu)⁶-Leu-Arg-Pro-(Azygly)NH₂
- v) deslorelin (US Patent 4 659 695)
 (pyr)Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-NH-CH₂-CH₂-NH₂
- vi) histerelin (US Patent 4 244 946)
 20 (pyr)Glu-His-Trp-Ser-Tyr-D-His(Bzl)-Leu-Arg-Pro-NH-CH₂-CH₃
- vii) avorelin (US 5 668 254)
 (pyr)Glu-His-Trp-Ser-Tyr-D-Trp(2-Me)-Leu-Arg-Pro-NH-CH₂-CH₃
- viii) nafarelin (US Patent 4 234 571)
 (pyr)Glu-His-Trp-Ser-Tyr-D-Nal(2)-Leu-Arg-Pro-NH-CH₂-CH₃;
- 25 lutrelin, cystorelin, gonadorelin or detirelix.

Particularly the LHRH agonist is selected from leuprorelin, buserelin, triptorelin and goserelin. More particularly the LHRH agonist is goserelin.

- Examples of suitable LHRH antagonists include, but are not limited to, antide, abarelix, antarelix, cetrorelix, azaline, ganirelix and those disclosed in US Patents 5 470 947
 30 (Folkers); 5 413 990 and 5 300 492 (Haviv); 5 371 070 (Koerber); 5 296 468 (Hoeger); 5 171 635 (Janaky); 5 003 011 and 4 431 635 (Coy); 4 992 421 (De); 4 801 577 (Nestor); and 4 851 385, 4 689 396 and 5 843 901 (Roeske).

- 7 -

Further examples of suitable LHRH antagonists include, but are not limited to the compounds described in WO 02/066477, WO 02/066478, WO 02/066459, WO 02/092565, PCT/GB03/003603 and PCT/GB03/003606, and the compounds described in these applications, particularly the compounds of claim 1 and the named examples are incorporated
5 herein by reference.

Particular bisphosphonates for use in the present invention are selected from tiludronic acid, ibandronic acid, incadronic acid, risedronic acid, zoledronic acid, clodronic acid, neridronic acid, pamidronic acid, alendronic acid, minodronic acid, olpadronic acid, TRK 530, CGP 47072, calcium clodronate or EB 1053. Further particular bisphosphonates for use
10 in the present invention are selected from etidronic acid, PNU-91638, NE-21650, NE-58025, NE-10790 or NE-10446.

Particular combinations of the present invention include:

- Compound (I) and leuprorelin, or a pharmaceutically acceptable salt thereof;
- Compound (I) and goserelin, or a pharmaceutically acceptable salt thereof;
- 15 • Compound (I) and abarelix, or a pharmaceutically acceptable salt thereof;
- Compound (I) and cetorelix, or a pharmaceutically acceptable salt thereof;
- Compound (I) and risedronic acid, or a pharmaceutically acceptable salt thereof;
- Compound (I) and zoledronic acid, or a pharmaceutically acceptable salt thereof;
- Compound (I) and clodronic acid, or a pharmaceutically acceptable salt thereof; and
- 20 • Compound (I) and pamidronic acid, or a pharmaceutically acceptable salt thereof.
- Compound (I) and alendronic acid, or a pharmaceutically acceptable salt thereof.

Suitable pharmaceutically-acceptable salts include, for example, salts with alkali metal (such as sodium, potassium or lithium), alkaline earth metals (such as calcium or magnesium), ammonium salts, and salts with organic bases affording physiologically acceptable cations,
25 such as salts with methylamine, dimethylamine, trimethylamine, piperidine and morpholine. In addition, for those compounds which are sufficiently basic, suitable pharmaceutically-acceptable salts include, pharmaceutically-acceptable acid-addition salts with hydrogen halides, sulphuric acid, phosphoric acid and with organic acids such as citric acid, maleic acid, methanesulphonic acid and p-toluenesulphonic acid. Alternatively, the
30 compounds may exist in zwitterionic form.

Therefore according to the present invention, there is provided a combination, comprising Compound (I) and an LHRH analogue for use as a medicament.

- 8 -

Therefore according to the present invention, there is provided a combination, comprising Compound (I) and a bisphosphonate for use as a medicament.

Therefore according to the present invention, there is provided a combination, comprising Compound (I), an LHRH analogue and a bisphosphonate for use as a
5 medicament.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I) and an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical
10 composition which comprises Compound (I) and a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I), an LHRH analogue and a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical
15 composition which comprises Compound (I), in association with a pharmaceutically acceptable diluent or carrier, in combination with a pharmaceutical composition which comprises an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical
20 composition which comprises Compound (I), in association with a pharmaceutically acceptable diluent or carrier, in combination with a pharmaceutical composition which comprises a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical
25 composition which comprises Compound (I), in association with a pharmaceutically acceptable diluent or carrier, in combination with a pharmaceutical composition which comprises an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier and a bisphosphonate in association with a pharmaceutically acceptable diluent or
30 carrier.

Therefore according to the present invention, there is provided a method of treating cancer, in a warm-blooded animal, such as man, in need of such treatment which comprises

- 9 -

administering to said animal an effective amount of Compound (I) in combination with an effective amount of an LHRH analogue.

Therefore according to the present invention, there is provided a method of treating cancer, in a warm-blooded animal, such as man, in need of such treatment which comprises
5 administering to said animal an effective amount of Compound (I) in combination with an effective amount of a bisphosphonate.

Therefore according to the present invention, there is provided a method of treating cancer, in a warm-blooded animal, such as man, in need of such treatment which comprises
10 administering to said animal an effective amount of Compound (I) in combination with an effective amount of an LHRH analogue and an effective amount of a bisphosphonate.

For the avoidance of doubt, where the treatment of cancer is indicated, it is to be understood that this also refers to the prevention of metastases and the treatment of metastases, i.e. cancer spread. Therefore the combination of the present invention could be used to treat a patient who has no metastases to stop them occurring, or to lengthen the time
15 period before they occur, and to a patient who already has metastases to treat the metastases themselves. Furthermore the treatment of cancer also refers to treatment of an established primary tumour or tumours and developing primary tumour or tumours. In one aspect of the invention the treatment of cancer relates to the prevention of metastases. In another aspect of the invention the treatment of cancer relates to the treatment of metastases. In another aspect
20 of the invention the treatment of cancer relates to treatment of an established primary tumour or tumours or developing primary tumour or tumours. Herein, the treatment of cancer also refers to the prevention of cancer *per se*.

In addition the treatment of cancer also refers to the production of an anti-angiogenic effect in a warm blooded animal.

25 According to a further aspect of the present invention there is provided a kit comprising Compound (I) and an LHRH analogue; optionally with instructions for use.

According to a further aspect of the present invention there is provided a kit comprising Compound (I), and a bisphosphonate; optionally with instructions for use.

According to a further aspect of the present invention there is provided a kit
30 comprising Compound (I), an LHRH analogue and a bisphosphonate; optionally with instructions for use.

According to a further aspect of the present invention there is provided a kit comprising:

- 10 -

- a) Compound (I), in a first unit dosage form;
- b) an LHRH analogue; in a second unit dosage form; and
- c) container means for containing said first and second dosage forms; and optionally
- d) with instructions for use.

5 An example of a unit dosage form for Compound (I) might be a tablet for oral formulation, see that described herein below. An example of a unit dosage form for an LHRH analogue might be a prolonged release formulation for an LHRH agonist (see herein below) or a prolonged release formulation or a tablet formulation for an LHRH antagonist (see herein below).

10 According to a further aspect of the present invention there is provided a kit comprising:

- a) Compound (I), in a first unit dosage form;
- b) a bisphosphonate; in a second unit dosage form; and
- c) container means for containing said first and second dosage forms; and optionally
- 15 d) with instructions for use.

 An example of a unit dosage form for a bisphosphonate might be a tablet for oral formulation, see herein below.

 According to a further aspect of the present invention there is provided a kit comprising:

- 20 a) Compound (I), in a first unit dosage form;
- b) an LHRH analogue; in a second unit dosage form; and
- c) a bisphosphonate; in a third unit dosage form; and
- d) container means for containing said first, second and third dosage forms; and optionally
- e) with instructions for use.

25 According to a further aspect of the present invention there is provided a kit comprising:

- a) Compound (I), together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) an LHRH analogue, in a second unit dosage form; and
- 30 c) container means for containing said first and second dosage forms; and optionally
- d) with instructions for use.

 According to a further aspect of the present invention there is provided a kit comprising:

- 11 -

- a) Compound (I), together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) a bisphosphonate, in a second unit dosage form; and
- c) container means for containing said first and second dosage forms; and optionally
- 5 d) with instructions for use.

According to a further aspect of the present invention there is provided a kit comprising:

- a) Compound (I), together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- 10 b) an LHRH analogue, in a second unit dosage form; and
- c) a bisphosphonate, in a third unit dosage form; and
- d) container means for containing said first, second and third dosage forms; and optionally
- e) with instructions for use.

According to a further aspect of the invention there is provided a pharmaceutical
15 composition which comprises Compound (I) and an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I) and a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.

20 According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I), an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier and a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.

25 According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I), in association with a pharmaceutically acceptable diluent or carrier, in combination with a pharmaceutical composition which comprises an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.

30 According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I), in association with a pharmaceutically acceptable diluent or carrier, in combination with a pharmaceutical composition which comprises a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.

- 12 -

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises Compound (I), in association with a pharmaceutically acceptable diluent or carrier, in combination with a pharmaceutical composition which comprises an LHRH analogue in association with a pharmaceutically acceptable diluent or carrier and a bisphosphonate in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.

The pharmaceutical compositions may be in a form suitable for oral administration, for example as a tablet or capsule, for parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion) as a sterile solution, suspension or emulsion, for topical administration as an ointment or cream or for rectal administration as a suppository. In general the above compositions may be prepared in a conventional manner using conventional excipients.

For example Compound (I) can be formulated as a tablet using the following excipients:

Compound (I);
Lactose monohydrate (filler);
Croscarmellose sodium (disintegrant);
Povidone (binder);
Magnesium stearate (lubricant);
Hypromellose (film coat component);
Polyethylene glycol 300 (film coat component); and
Titanium dioxide (film coat component).

According to a further aspect of the present invention there is provided a kit comprising Compound (I) and an LHRH analogue; optionally with instructions for use; for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising Compound (I), and a bisphosphonate; optionally with instructions for use; for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising Compound (I), an LHRH analogue and a bisphosphonate; optionally with instructions for use; for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising:

- 13 -

- a) Compound (I), in a first unit dosage form;
 - b) an LHRH analogue in a second unit dosage form; and
 - c) container means for containing said first and second dosage forms; and optionally
 - d) with instructions for use;
- 5 for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising:

- a) Compound (I), in a first unit dosage form;
 - b) a bisphosphonate in a second unit dosage form; and
 - 10 c) container means for containing said first and second dosage forms; and optionally
 - d) with instructions for use;
- for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising:

- 15 a) Compound (I), in a first unit dosage form;
 - b) an LHRH analogue in a second unit dosage form; and
 - c) a bisphosphonate in a third unit dosage form; and
 - d) container means for containing said first, second and third dosage forms; and optionally
 - e) with instructions for use;
- 20 for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising:

- a) Compound (I), together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
 - 25 b) an LHRH analogue in a second unit dosage form; and
 - c) container means for containing said first and second dosage forms; and optionally
 - d) with instructions for use;
- for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a kit comprising:

- 30 a) Compound (I), together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) a bisphosphonate in a second unit dosage form; and

- 14 -

- c) container means for containing said first and second dosage forms; and optionally
 - d) with instructions for use;
- for use in the treatment of cancer.

- According to a further aspect of the present invention there is provided a kit
- 5 comprising:
- a) Compound (I), together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
 - b) an LHRH analogue in a second unit dosage form; and
 - c) a bisphosphonate in a second unit dosage form; and
- 10 d) container means for containing said first, second and third dosage forms; and optionally
- e) with instructions for use;
- for use in the treatment of cancer.

- According to another feature of the invention there is provided the use of Compound (I), in combination with an LHRH analogue in the manufacture of a medicament for use in the
- 15 treatment of cancer, in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of Compound (I), in combination with a bisphosphonate in the manufacture of a medicament for use in the treatment of cancer, in a warm-blooded animal, such as man.

- According to another feature of the invention there is provided the use of Compound
- 20 (I), in combination with an LHRH analogue and a bisphosphonate in the manufacture of a medicament for use in the treatment of cancer, in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of Compound (I), in combination with an LHRH analogue in the treatment of cancer, in a warm-blooded animal, such as man.

- 25 According to another feature of the invention there is provided the use of Compound (I), in combination with a bisphosphonate in the treatment of cancer, in a warm-blooded animal, such as man.

- According to another feature of the invention there is provided the use of Compound (I), in combination with an LHRH analogue and a bisphosphonate in the treatment of cancer,
- 30 in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a combination comprising Compound (I) and an LHRH analogue for use in the treatment of cancer.

- 15 -

According to a further aspect of the present invention there is provided a combination comprising Compound (I), and a bisphosphonate for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a combination comprising Compound (I), an LHRH analogue and a bisphosphonate for use in the treatment
5 of cancer.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of Compound (I), optionally together with a pharmaceutically acceptable diluent or carrier, in combination with an effective amount of an LHRH analogue optionally together with a pharmaceutically
10 acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of Compound (I), optionally together with a pharmaceutically acceptable diluent or carrier, in combination with an
15 effective amount of a bisphosphonate optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment for use in the treatment of cancer.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of Compound (I), optionally
20 together with a pharmaceutically acceptable diluent or carrier, in combination with an effective amount of an LHRH analogue optionally together with a pharmaceutically acceptable diluent or carrier and an effective amount of a bisphosphonate optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment for use in the treatment of cancer.

25 The amount of Compound (I), or a pharmaceutically acceptable salt thereof, administered would be that sufficient to provide the desired pharmaceutical effect. For instance, Compound (I) could be administered to a warm-blooded animal orally, at a unit dose less than 1g daily but more than 2.5mg. Particularly Compound (I) could be administered to a warm-blooded animal, at a unit dose of less than 250 mg per day. In another aspect of the
30 invention, Compound (I) could be administered to a warm-blooded animal, at a unit dose of less than 130 mg per day. In a further aspect of the invention, Compound (I) could be administered to a warm-blooded animal, at a unit dose of less than 50 mg per day.

- 16 -

LHRH analogues, particularly LHRH agonists would normally be administered as a prolonged release formulation in order to enhance the therapeutic effectiveness of the drug. A number of prolonged release formulations containing LHRH analogues are known and can be achieved by administering the drug in the form of a biodegradable polymer matrix which releases the drug over a prolonged period. Suitable biodegradable polymers are polylactides. The term 'polylactide' is used in a generic sense to include polymers of lactic acid alone, copolymers of lactic and glycolic acid, mixtures of such polymers, mixtures of such copolymers and mixtures of such polymers and copolymers, the lactic acid being either in racemic or optically active form.

One approach has been to develop microparticle or microcapsule formulations comprising the biodegradable polymer, typically a poly(lactide-co-glycolide) co-polymer, in which the LHRH analogue is microencapsulated.

EP 839 525 describes the preparation of microcapsules containing LHRH analogues using an emulsion based process wherein a water-in-oil emulsion is prepared comprising an inner aqueous phase containing the LHRH analogue and an outer oil phase comprising a solution of a polymer of lactic acid in an organic solvent. Generally, microencapsulated LHRH analogues are suitable for delivering the LHRH analogue for up to 3 months.

EP 058 481 describes monolithic implants comprising a biodegradable poly(lactide-co-glycolide) co-polymer and an LHRH agonist.

WO 03/022297 describes monolithic implants prepared using a combination of a specific polylactide polymer and a LHRH analogue which continuously releases the LHRH analogue over a period of at least six months when placed in an aqueous physiological-type environment.

WO 03/022243 describes the sustained release of microcrystalline peptide suspensions containing LHRH agonists.

As an alternative Eligard (an LHRH agonist) is sold as a flowable polymeric formulation composed of poly (DL-lactide) which is suspended in a non-aqueous solvent N-methyl-2-pyrrolidone.

An LHRH antagonist as well as being delivered by any of the prolonged release methods listed above could also be prepared for oral administration using standard techniques and known excipients.

The LHRH analogue will normally be administered to a warm-blooded animal at a unit dose, for example, from about 3-4mg per month, or 10-11mg every 3 months, of active

- 17 -

ingredient. When the LHRH analogue is goserelin, a sustained release subcutaneous formulation containing 3.6mg per month, or 10.6mg every 3 months, of active ingredient. Conveniently the daily oral dose of leuprorelin is 3.75mg per month, or 11.25mg every 3 months. The optimum dosage however, may be determined by the practitioner who is treating
5 any particular patient.

The bisphosphonate will normally be administered to a warm-blooded animal at a unit dose, of an amount known to the skilled practitioner as a therapeutically effective dose. For a single dosage form, the active ingredients may be compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight
10 of the total composition. Dosage unit forms will generally contain about 20 mg to about 500 mg of each active ingredient. However the daily dose will necessarily be varied depending upon the host treated, the particular route of administration, and the severity of the illness being treated. Accordingly the optimum dosage may be determined by the practitioner who is treating any particular patient.

15 The dosage of each of the drugs and their proportions have to be composed so that the best possible treatment effects, as defined by national and international guidelines (which are periodically reviewed and re-defined), will be met.

- 18 -

Claims

1. A combination, comprising N-(3-methoxy-5-methylpyrazin-2-yl)-2-(4-[1,3,4-oxadiazol-2-yl]phenyl)pyridine-3-sulphonamide, or a pharmaceutically acceptable salt thereof, and an LHRH analogue.
5
2. A combination, comprising N-(3-methoxy-5-methylpyrazin-2-yl)-2-(4-[1,3,4-oxadiazol-2-yl]phenyl)pyridine-3-sulphonamide, or a pharmaceutically acceptable salt thereof, and a bisphosphonate.
10
3. A combination, comprising N-(3-methoxy-5-methylpyrazin-2-yl)-2-(4-[1,3,4-oxadiazol-2-yl]phenyl)pyridine-3-sulphonamide, or a pharmaceutically acceptable salt thereof, an LHRH analogue and a bisphosphonate.
15
4. A combination according to claims 1 or 3 wherein the LHRH analogue is an LHRH agonist.
20
5. A combination according to claim 4 wherein the LHRH agonist is selected from leuporelin, buserelin, triptorelin and goserelin.
25
6. A combination according to claims 4 or 5 wherein the LHRH agonist is goserelin.
7. A combination according to claims 1 or 3 wherein the LHRH analogue is an LHRH antagonist.
30
8. A combination according to claim 7 wherein the LHRH antagonist is selected from antide, abarelix, antarelix, cetrotorelix, azaline or ganirelix.
9. A combination according to claims 2 - 8 wherein the bisphosphonate is selected from tiludronic acid, ibandronic acid, incadronic acid, risedronic acid, zoledronic acid, clodronic acid, neridronic acid, pamidronic acid, alendronic acid, minodronic acid, olpadronic acid, TRK 530, CGP 47072, calcium clodronate or EB 1053.

- 19 -

10. A combination as claimed in any one of claims 1-9 for use as a medicament.
11. A pharmaceutical composition as claimed in any one of claims 1-9 in association with a pharmaceutically acceptable diluent or carrier.
- 5 12. A method of treating cancer, in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a combination as claimed in any one of claims 1-9.
- 10 13. A pharmaceutical composition which comprises a combination as claimed in any one of claims 1-9 in association with a pharmaceutically acceptable diluent or carrier for use in the treatment of cancer.
14. The use of a combination as claimed in any one of claims 1-9 in the manufacture of a
15 medicament for use in the treatment of cancer, in a warm-blooded animal, such as man.

INTERNATIONAL SEARCH REPORT

PCT/GB2004/003733

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K31/497 A61K31/663 A61K31/675 A61K38/09 A61P35/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, EMBASE, BIOSIS, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	DATABASE WPI Section Ch, Week 200329 Derwent Publications Ltd., London, GB; Class B05, AN 2003-300573 XP002309704 & WO 03/015820 A1 (TAKEDA CHEM IND LTD) 27 February 2003 (2003-02-27) abstract	1-14
P, Y	& EP 1 424 080 A (TAKEDA CHEMICAL INDUSTRIES LTD) 2 June 2004 (2004-06-02) page 17, line 44, paragraph 12 - page 19, line 55; claims 21,22,31-33	1-14
Y	US 2003/092757 A1 (SINGH AMITABH ET AL) 15 May 2003 (2003-05-15) paragraphs '0024!, '0173!; claim 1 ----- -/-	1-14

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the International filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the International filing date but later than the priority date claimed

T later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the International search

9 December 2004

Date of mailing of the International search report

30/12/2004

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Blott, C

INTERNATIONAL SEARCH REPORT

PCT/GB2004/003733

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	"ZD4054: a specific endothelin A receptor antagonist with potential utility in prostate cancer and metastatic bone disease" EUROPEAN JOURNAL OF CANCER, PERGAMON PRESS, OXFORD, GB, vol. 38, November 2002 (2002-11), page S102, XP004403782 ISSN: 0959-8049 the whole document	1-14
Y	WO 02/087555 A (NOVARTIS ERFINDE VERWALT GMBH ; NOVARTIS AG (CH); SEAMAN JOHN J (US)) 7 November 2002 (2002-11-07) page 14, lines 9-15	1-14
A	MORRIS MICHAEL J ET AL: "Clinical approaches to osseous metastases in prostate cancer." THE ONCOLOGIST. 2003, vol. 8, no. 2, 2003, pages 161-173, XP002309703 ISSN: 1083-7159 page 164, column 2 page 168, column 2; table 3	1-14
P,X	WO 2004/035057 A (HANCOX URSULA JOY ; TONGE DAVID WILLIAM (GB); CURWEN JON OWEN (GB); JO) 29 April 2004 (2004-04-29) page 8, line 11 - page 10, line 13; claims 4,12	1-14
P,X	WO 2004/032922 A (CURWEN JON OWEN ; JOHNSTONE DONNA (GB); ASTRAZENECA UK LTD (GB); HUGHE) 22 April 2004 (2004-04-22) page 8, line 18 - page 10, line 21 claims 11,14,17	1-14

INTERNATIONAL SEARCH REPORT

PCT/GB2004/003733

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 03015820	A1	27-02-2003	CA	2458452 A1	27-02-2003
			EP	1424080 A1	02-06-2004
			JP	2003137814 A	14-05-2003
EP 1424080	A	02-06-2004	CA	2458452 A1	27-02-2003
			EP	1424080 A1	02-06-2004
			WO	03015820 A1	27-02-2003
			JP	2003137814 A	14-05-2003
US 2003092757	A1	15-05-2003	NONE		
WO 02087555	A	07-11-2002	BR	0209365 A	08-06-2004
			CA	2443625 A1	07-11-2002
			CZ	20032950 A3	16-06-2004
			WO	02087555 A2	07-11-2002
			EP	1418897 A2	19-05-2004
			HU	0400096 A2	28-04-2004
			JP	2004528340 T	16-09-2004
			NO	20034877 A	19-12-2003
			SK	13512003 A3	03-08-2004
			US	2004157799 A1	12-08-2004
			ZA	200307666 A	05-05-2004
WO 2004035057	A	29-04-2004	WO	2004035057 A1	29-04-2004
WO 2004032922	A	22-04-2004	WO	2004032922 A1	22-04-2004